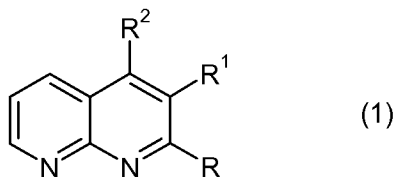


AMENDMENTS TO THE CLAIMS

1. (Currently amended) The compound of the general formula (1):



wherein

R is halo;

R¹ is aryl ~~or~~ heteroaryl; ~~morpholino, piperidino or pyrrolidino;~~

R² is NR³R⁴,

wherein R³ and R⁴ are independently H, C₁₋₈ alkyl, C₂₋₈ alkenyl, C₂₋₈ alkynyl, ~~aryl, aryl(C₁₋₈)alkyl, C₃₋₈ cycloalkyl, C₃₋₈ cycloalkyl(C₁₋₈)alkyl, heteroaryl, heteroaryl(C₁₋₈)alkyl, NR⁵R⁶, provided that not both R³ and R⁴ are H or NR⁵R⁶;~~

or wherein R³ and R⁴ together form a C₃₋₇ alkylene or C₃₋₇ alkenylene chain optionally substituted with one or more C₁₋₄ alkyl or C₁₋₄ alkoxy groups;

or wherein R³ and R⁴ together with the nitrogen atom to which they are attached form a morpholine, thiomorpholine, thiomorpholine S-oxide or thiomorpholine S-dioxide ring or a piperazine or piperazine N-(C₁₋₄)alkyl (especially N-methyl) ring;

~~R⁵ and R⁶ are independently H, C₁₋₈ alkyl, C₂₋₈ alkenyl, C₂₋₈ alkynyl, aryl, aryl(C₁₋₈)alkyl, C₃₋₈ cycloalkyl, C₃₋₈ cycloalkyl(C₁₋₈)alkyl, heteroaryl or hetero-aryl(C₁₋₈)alkyl;~~

and wherein

said alkyl, alkenyl, ~~or~~ alkynyl ~~or cycloalkyl~~ groups or moieties are optionally substituted with halogen, cyano, C₁₋₆alkoxy, C₁₋₆alkylcarbonyl, C₁₋₆alkoxycarbonyl, C₁₋₆haloalkoxy, C₁₋₆alkylthio, tri(C₁₋₄)alkylsilyl, C₁₋₆alkylamino or C₁₋₆dialkylamino;

said morpholine, thiomorpholine, ~~piperidine, and~~ piperazine ~~and pyrrolidino~~ rings are optionally substituted with C₁₋₄ alkyl (especially methyl); and

said aryl or heteroaryl groups or moieties are optionally substituted with one or more substituents selected from the group consisting halo, hydroxy, mercapto, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, C₁₋₆alkoxy, C₂₋₆alkenyloxy, C₂₋₆alkynyloxy, halo(C₁₋₆)alkyl, halo(C₁₋₆)alkoxy, C₁₋₆alkylthio, halo(C₁₋₆)alkylthio, hydroxy(C₁₋₆)alkyl, C₁₋₄alkoxy(C₁₋₆)alkyl, C₃₋₆cycloalkyl, C₃₋₆cycloalkyl(C₁₋₄)alkyl, phenoxy, benzyloxy, benzoyloxy, cyano, isocyano, thiocyanato, isothiocyanato, nitro, -NR^{'''}R^{'''}, -NHCOR^{'''}, -NHCONR^{'''}R^{'''}, -CONR^{'''}R^{'''}, -SO₂R^{'''}, -OSO₂R^{'''},

-COR''', -CR'''=NR''' and -N=CR'''R''', in which R''' and R''' are independently hydrogen, C₁₋₄ alkyl, halo(C₁₋₄)alkyl, C₁₋₄ alkoxy, halo(C₁₋₄)alkoxy, C₁₋₄ alkylthio, C₃₋₆ cycloalkyl, C₃₋₆ cycloalkyl(C₁₋₄) alkyl, phenyl or benzyl, the phenyl and benzyl groups being optionally substituted with halogen, C₁₋₄ alkyl or C₁₋₄ alkoxy.

2. (Currently amended) A compound according claim 1 wherein:

(A) R³ is C₁₋₈ alkyl, halo(C₁₋₈) alkyl, hydroxy(C₁₋₈)alkyl, C₁₋₄ alkoxy(C₁₋₈)alkyl, C₁₋₄ alkoxyhalo(C₁₋₈)alkyl, tri(C₁₋₄)alkylsilyl(C₁₋₆)alkyl, C₁₋₄ alkylcarbonyl(C₁₋₈)alkyl, C₁₋₄ alkylcarbonylhalo(C₁₋₈)alkyl, phenyl(₁₋₄) alkyl, C₂₋₈ alkenyl, halo(C₂₋₈)alkenyl, C₂₋₈ alkynyl, ~~C₃₋₈ cycloalkyl optionally substituted with chloro, fluoro or methyl, C₃₋₈ cycloalkyl (C₁₋₄) alkyl, phenylamino, piperidino or morpholino, the phenyl ring of phenylalkyl or phenylamino being optionally substituted with one, two or three substituents selected from halo, C₁₋₄ alkyl, halo(C₁₋₄)alkyl, C₁₋₄ alkoxy and halo (C₁₋₄)alkoxy; and R⁴ is H, C₁₋₄ alkyl, halo(C₁₋₄)alkyl or amino; or~~

(B) R³ and R⁴ together form a C₃₋₇ alkylene or C₃₋₇ alkenylene chain optionally substituted with methyl; or

(C) R³ and R⁴, together with the nitrogen atom to which they are attached, form a morpholine, thiomorpholine, ~~thiomorpholine S-oxide or thiomorpholine S-dioxide ring or a piperazine or piperazine N-(C₁₋₄)alkyl (especially N-methyl) ring, in which the morpholine or piperazine rings are optionally substituted with methyl.~~

3. (Currently amended) A compound according to claim1 wherein R¹ is phenyl optionally substituted with from one to five halogen atoms or with from one to three substituents selected from halo, C₁₋₄ alkyl, halo(C₁₋₄)alkyl, C₁₋₄ alkoxy or halo(C₁₋₄)alkoxy, ~~pyridyl optionally substituted with from one to four halogen atoms or with from one to three substituents selected from halo, C₁₋₄ alkyl, halo(C₁₋₄)alkyl, C₁₋₄ alkoxy or halo(C₁₋₄)alkoxy, 2- or 3-thienyl optionally substituted with from one to three halogen atoms or with from one to three substituents selected from halo, C₁₋₄ alkyl, halo(C₁₋₄)alkyl, C₁₋₄ alkoxy or halo(C₁₋₄)alkoxy, or piperidino or morpholino both optionally substituted with one or two methyl groups.~~

4. (Original) A compound according to claim 3 wherein R¹ is 2,6-difluorophenyl, 2-fluoro-6-chlorophenyl, 2,5,6-trifluorophenyl, 2,4,6-trifluorophenyl, 2,6-difluoro-4-methoxyphenyl or pentafluorophenyl.

5. Cancelled.

6. (Currently amended) A compound according to claim 1 wherein:

(A) R^3 is ~~C_{1-4} alkyl, C_{1-8} alkyl, halo(C_{1-4})alkyl, C_{2-4} alkenyl, C_{3-6} cycloalkyl, C_{3-6} cycloalkyl(C_{1-4})alkyl or phenylamino in which the phenyl ring is optionally substituted with one, two or three substituents selected from halo, C_{1-4} alkyl, halo(C_{1-4})alkyl, C_{1-4} alkoxy and halo(C_{1-4})alkoxy, and R^4 is H, or C_{1-4} alkyl, or amine;~~

(B) or wherein R^3 and R^4 together form a ~~C_{4-6} C_{3-7}~~ alkylene chain optionally substituted with C_{1-4} alkyl, or C_{1-4} alkoxy;

(C) or wherein R^3 and R^4 , together with the nitrogen atom to which they are attached, form a morpholine, ~~thiomorpholine, thiomorpholine S-oxide or thiomorpholine S-dioxide ring or a~~ piperazine or piperazine N -(C_{1-4})alkyl (especially N -methyl) ring; and

wherein said alkyl, ~~or, alkenyl, alkynyl or cycloalkyl~~ groups or moieties are optionally substituted with halogen, cyano, C_{1-6} alkoxy, C_{1-6} alkylcarbonyl, C_{1-6} alkoxycarbonyl, C_{1-6} haloalkoxy, C_{1-6} alkylthio, tri(C_{1-4})alkylsilyl, C_{1-6} alkylamino or C_{1-6} dialkylamino;

and wherein said ~~said morpholine and, thiomorpholine, piperidine, piperazine and pyrrolidine~~ rings are optionally substituted with C_{1-4} alkyl;

and wherein said aryl or heteroaryl groups or moieties are optionally substituted with one or more substituents selected from the group consisting of halo, hydroxy, mercapto, C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, C_{1-6} alkoxy, C_{2-6} alkenyloxy, C_{2-6} alkynyloxy, halo(C_{1-6})alkyl, halo(C_{1-6})alkoxy, C_{1-6} alkylthio, halo(C_{1-6})alkylthio, hydroxy(C_{1-6})alkyl, C_{1-4} alkoxy(C_{1-6})alkyl, C_{3-6} cycloalkyl, C_{3-6} cycloalkyl(C_{1-4})alkyl, phenoxy, benzyloxy, benzoyloxy, cyano, isocyano, thiocyanato, isothiocyanato, nitro, $-NR''R'''$, $-NHCOR''$, $-NHCONR''R'''$, $-CONR''R'''$, $-SO_2R''$, $-OSO_2R''$, $-COR''$, $-CR''=NR'''$ and $-N=CR''R'''$, in which R'' and R''' are independently hydrogen, C_{1-4} alkyl, halo(C_{1-4})alkyl, C_{1-4} alkoxy, halo(C_{1-4})alkoxy, C_{1-4} alkylthio, C_{3-6} cycloalkyl, C_{3-6} cycloalkyl(C_{1-4})alkyl, phenyl or benzyl, the phenyl and benzyl groups being optionally substituted with halogen, C_{1-4} alkyl or C_{1-4} alkoxy.

7. (Previously presented) A compound according to claim 1 wherein R^1 is optionally substituted phenyl.

8. (Currently amended) A compound according to claim 1 wherein:

R^1 is phenyl optionally substituted with from one to five halogen atoms or with from one to three substituents selected from the group consisting of halo, C_{1-4} alkyl, halo(C_{1-4})alkyl, C_{1-4} alkoxy and ~~or halo(C_{1-4})alkoxy, and pyridyl optionally substituted with from one to four halogen atoms or with from one to three substituents selected from halo, C_{1-4} alkyl, halo(C_{1-4})alkyl, C_{1-4} alkoxy or~~

~~halo(C₁₋₄)alkoxy, 2- or 3-thienyl optionally substituted with from one to three halogen atoms or with from one to three substituents selected from halo, C₁₋₄alkyl, halo(C₁₋₄)alkyl, C₁₋₄alkoxy or halo(C₁₋₄)alkoxy, or piperidino or morpholino both optionally substituted with one or two methyl groups; and~~

wherein R³ is ~~C₁₋₄ alkyl or halo(C₁₋₄) alkyl; C₁₋₆ alkyl, halo(C₁₋₆)alkyl, hydroxy(C₁₋₆)alkyl, C₁₋₄ alkoxy(C₁₋₆)alkyl, C₁₋₄ alkoxyhalo(C₁₋₆)alkyl, tri(C₁₋₄)alkylsilyl(C₁₋₆)alkyl, C₁₋₄ alkylcarbonyl(C₁₋₆)alkyl, C₁₋₄ alkylcarbonylhalo(C₁₋₆)alkyl, phenyl(C₁₋₄)alkyl, C₂₋₈ alkenyl, halo(C₂₋₈)alkenyl, C₂₋₈ alkynyl, C₂₋₈ cycloalkyl optionally substituted with chloro, fluoro or methyl, C₂₋₈ cycloalkyl(C₁₋₄)alkyl, phenylamino, piperidino or morpholino, the phenyl ring of phenylalkyl or phenylamino being optionally substituted with one, two or three substituents selected from halo, C₁₋₄ alkyl, halo(C₁₋₄)alkyl, C₁₋₄ alkoxy and halo(C₁₋₄)alkoxy; and R⁴ is H, C₁₋₄ alkyl, halo(C₁₋₄)alkyl or amino;~~

or wherein R³ and R⁴ together form a ~~C₁₋₆ alkylene chain, C₃₋₇ alkylene or C₃₋₇ alkenylene chain optionally substituted with methyl;~~

~~or wherein, or, R³ and R⁴ together with the nitrogen atom to which they are attached, R³ and R⁴ form a morpholine, thiomorpholine, thiomorpholine S-oxide or thiomorpholine S-dioxide ring or a piperazine or piperazine N-(C₁₋₄)alkyl (especially N-methyl) ring, in which the morpholine or piperazine rings are optionally substituted with methyl.~~

9. (Currently amended) A compound according to claim 1 wherein:

R¹ is phenyl optionally substituted with from one to five halogen atoms; ~~or with from one to three substituents selected from halo, C₁₋₄ alkyl, halo(C₁₋₄)alkyl, C₁₋₄alkoxy or halo(C₁₋₄)alkoxy; and~~

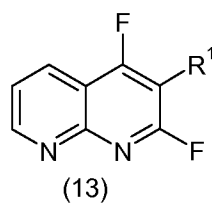
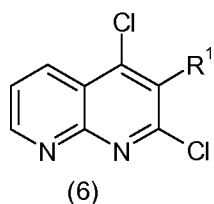
wherein R³ is ~~C₁₋₄ alkyl, halo(C₁₋₄)alkyl, C₂₋₄alkenyl, C₂₋₆ cycloalkyl, C₃₋₆ cycloalkyl(C₁₋₄)alkyl or phenylamino in which the phenyl ring is optionally substituted with one, two or three substituents selected from halo, C₁₋₄ alkyl, halo(C₁₋₄)alkyl, C₁₋₄ alkoxy and halo(C₁₋₄)alkoxy; and R⁴ is H, C₁₋₄ alkyl or amino;~~

or wherein R³ and R⁴ together form a C₄₋₆ alkylene chain optionally substituted with methyl;

or wherein R³ and R⁴, together with the nitrogen atom to which they are attached, form a morpholine ring.

10. (Previously presented) A process for preparing a compound of the general formula (1) according to claim 1 wherein R is chloro or fluoro, comprising:

(A) reacting an amine of the general formula NR³R⁴ with a compound of the general formula (6) or (13):



wherein R¹, R³ and R⁴ are as defined in claim 1.

11. (Original): A plant fungicidal composition comprising a fungicidally effective amount of a compound as defined in claim 1 and a suitable carrier or diluent therefor.

12. (Previously presented) A method of combating or controlling phytopathogenic fungi which comprises applying to a plant, to a seed of a plant, to the locus of the plant or seed or to soil or to any other plant growth medium, a fungicidally effective amount of a compound according to claim 1.